

Compound odontoma with cystic proliferation. a case report

1 Rojas Ávila Josué Julián*Alonso Moctezuma Alejandro * Díaz Ayala Francisco*

1 First resident oral and Maxillofacial Surgery Universidad Nacional Autónoma de México, México 1Chairman of Oral and Maxillofacial Surgery Universidad Nacional Autónoma de México, México, *Chief Resident Oral and Maxillofacial Surgery, Universidad Nacional Autónoma de México, México

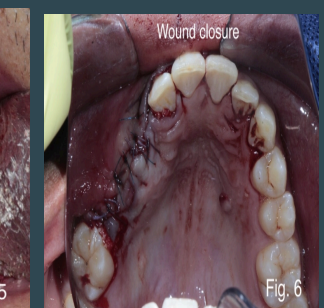
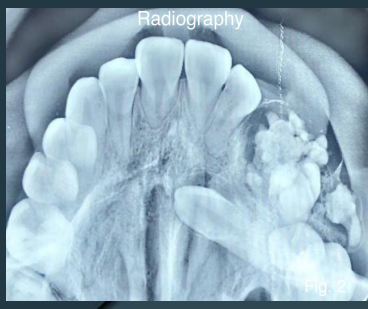
INTRODUCTION

Odontoma is a non aggressive ectomesenchymal tumor of unknown origin that are more considered developmental malformations (hamartomas) than benign neoplasms. They are typically diagnosed during the first two decades of life and have no sex predilection. The etiology is unknown, but genetic mutation in a tooth germ is a possible factor^{1,2}.

OBJECTIVE

Describe the clinical and management of a compound odontoma with cystic proliferation in a case report.

CASE REPORT A 24 years old patient who reports increased volume in the upper left vestibular region (fig.1) with one year evolution and pain, without sistemic diseases. Radiografically was observed a mixed image well definated, causing retention og o.d 23 and 25(fig.2) it was decided to perform enucleation and regeneration with allograft (fig. 5) and diagnosticated like compound odontoma with cystic proliferationin (fig.4) at the clinical for Oral and Maxillofacial Surgery of the Universidad Nacional Autónoma de México.



Histopathology: Consist chiefly of enamel and dentin, with variable amounts of pulp and cementum and cystic proliferation with a final diagnosis of cystic odontoma after two years of evolution without recurrence.

DISCUSSION:

Clinically odontomas are either complex or compound and are classified as: intraosseous: odontomas occurring inside the bone and may erupt (erupted odontomas) into the oral cavity and extraosseous: odontomas occurring in the soft tissue overing the tooth bearing portion of the jaws¹⁻³. Astekar suggested Complete surgical excision under local anaesthesia, Barnes et al. Suggested an enucleation^{2,3}. As a result of their odontogenic nature, including epithelial and mesenchymal tissues, odontomas can develop cystic transformation into a dentigerous cyst. This cyst results from the cystic degeneration of the enamel organ after partial or total development of the crown. These changes are cited as possible, but are rarely seen in clinical practice¹⁻⁴.in our case these same characteristics were observed

CONCLUSION:

The treatment for this entity must be the enucleation of the lesion. The association of an odontoma with cystic lesions are rare and should be considered to plan an adequate treatment of these lesions.

CONFLICTS OF INTEREST The authors declare no conflicts of interest

REFERENCES:

1. Ide F, Kikuchi K, Miyazaki Y, Kusama K. Archegonous Cystic Odontoma Is Not Necessarily Primordial. Head Neck Pathol. 2016;10(3):418-21.
2. Costa V, Carls AR, León JE, Ramos CJ, Jardini V, Kaminagakura E. Case Report Cystic Odontoma in a Patient with Hodgkin's Lymphoma. 2015;2015:5-10.
3. Astekar M, Manjunatha BS, Kaur P, Singh J. Histopathological insight of complex odontoma associated with adenigerous cyst. BMJ Case Rep 2014; 2014.
4. Barnes L, Eveson JW, Reichart P, Sidransky D. Odontogenic tumors. World Health Organization Classification of Tumours. Pathology and Genetics of Head and Neck Tumours. Lyon: IARCPress; 2017.